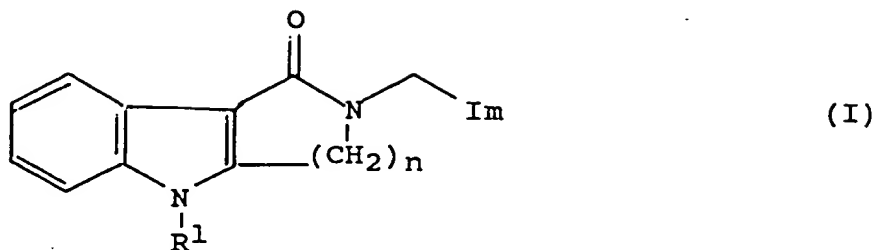
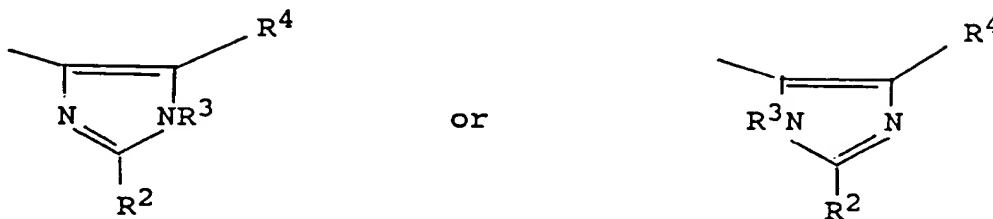


WE CLAIM:

1. A compound of formula (I)



wherein Im represents an imidazolyl group of the formula:



and R¹ represents a hydrogen atom or a group selected from C₁-6alkyl, C₃-6alkenyl, C₃-10alkynyl, C₃-7cycloalkyl, C₃-7cycloalkylC₁-4alkyl, phenyl, phenylC₁-3alkyl, phenylmethoxymethyl, phenoxyethyl, phoxymethyl; ~~-CO₂R⁵, -COR⁵, -CONR⁵R⁶ or -SO₂R⁵ (wherein R⁵ and R⁶, which may be the same or different, each represents a hydrogen atom, a C₁-6alkyl or C₃-7cycloalkyl group or a phenyl or phenylC₁-4alkyl group, in which the phenyl group is optionally substituted by one or more C₁-4alkyl, C₁-4alkoxy or hydroxy groups or halogen atoms, with the proviso that R⁵ does not represent a hydrogen atom when R¹ represents a group -CO₂R⁵ or -SO₂R⁵)~~

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27R
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one of the groups represented by R², R³ and R⁴ is a hydrogen atom or a C₁-6alkyl, C₃-7cycloalkyl, C₃-6alkenyl, phenyl or phenylC₁-3alkyl group, and each of the other two

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groups, which may be the same or different, represents a hydrogen atom or a C₁₋₆alkyl group;

n represents 2 or 3;

or a physiologically acceptable salt or solvate thereof.

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2. A compound according to claim 1 in which R¹ represents a C₁₋₄alkyl, C₃₋₄alkynyl, C₅₋₆cycloalkyl,

C C₅₋₆cycloalkylmethyl, phenylC₁₋₂alkyl, ^{or} phenylmethoxymethyl,
 C ~~or N,N-diC₁₋₃alkylcarboxamido group~~

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3. A compound according to claim 1 in which R², R³ and R⁴ each independently represent a hydrogen atom or a C₁₋₃alkyl group.

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4. A compound according to claim 1 in which R¹ represents a hydrogen atom or a C₁₋₄alkyl, C₃₋₄alkenyl, C₃₋₄alkynyl, C₅₋₆cycloalkyl, C₅₋₆cycloalkylmethyl,
 C phenylC₁₋₂alkyl, ^{or} phenylmethoxymethyl,
 C ~~N,N-diC₁₋₃alkylcarboxamido or C₁₋₃alkylsulphonyl group~~

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R² represents a hydrogen atom; and R³ and R⁴ each represent a hydrogen atom or a C₁₋₃alkyl group.

5. A compound according to claim 1 in which R¹ represents a methyl, n-propyl, prop-2-ynyl, cyclopentyl,
 C 25 cyclopentylmethyl, ^{or} benzyl ~~or N,N-dimethylcarboxamido group~~;
 R² and R³ each represent a hydrogen atom; and R⁴ represents a methyl group.

6. A compound according to claim 4 in which n represents 2.

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7. A compound according to claim 5 in which n represents 2.

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8. 2,3,4,5-Tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one;

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ona

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salt or solvate

and physiologically acceptable ~~salts and solvates~~ thereof.

9. A compound selected from:

2,3,4,5-Tetrahydro-5-(phenylmethyl)-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one;

5-cyclopentyl-2,3,4,5-tetrahydro-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one;

2,3,4,5-tetrahydro-2-[(5-methyl-1H-imidazol-4-yl)methyl]-5-propyl-1H-pyrido[4,3-b]indol-1-one;

5-(cyclopentylmethyl)-2,3,4,5-tetrahydro-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one;

3,4,5,6-tetrahydro-6-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-azepino[4,3-b]indol-1(2H)-one;

2,3,4,5-tetrahydro-N,N-dimethyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1-oxo-5H-pyrido[4,3-b]indole-5-carboxamide;

2,3,4,5-tetrahydro-2-[(5-methyl-1H-imidazol-4-yl)methyl-5-(2-propynyl)-1H-pyrido[4,3-b]indol-1-one;

and physiologically acceptable salts and solvates thereof.

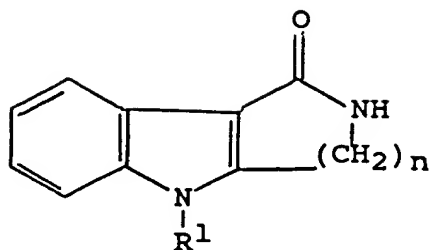
10. A compound according to claim 1 in the form of a hydrochloride, hydrobromide, sulphate, alkylsulphonate, arylsulphonate, phosphate, acetate, citrate, succinate, tartrate, fumarate or maleate salt.

11. The compound of claim 8 in the form of a hydrochloride salt.

12. The compound of claim 7 in the form of a maleate salt.

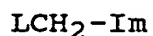
13. A process for the preparation of a compound of formula (I) as defined in claim 1 or a physiologically acceptable salt or solvate thereof, which process comprises:

(A) alkylating a compound of formula (II)



(II)

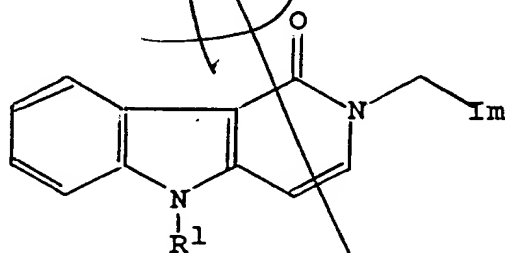
with a compound of formula (III)



(III)

or a protected derivative thereof, wherein L represents a leaving atom or group, followed if necessary by removal of any protecting groups present; or

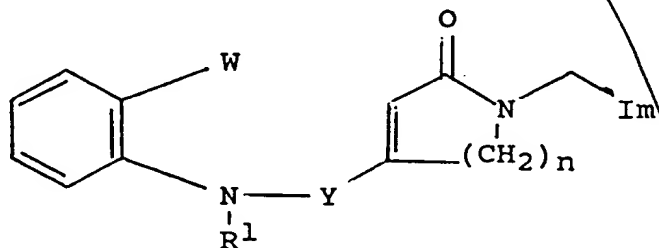
(B) for the preparation of a compound of formula (I) in which n is 2, hydrogenating a compound of formula (IV)



(IV)

or a protected derivative thereof, followed if necessary by removal of any protecting groups present; or

(C) cyclising a compound of formula (V)



(V)

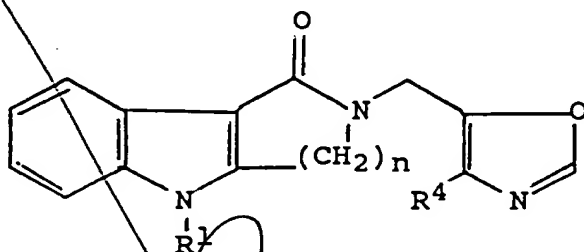
wherein W represents a hydrogen atom and Y represents the group NH, or W represents a halogen atom and Y represents a bond, or a salt or protected derivative thereof, followed if necessary by removal of any protecting groups present;

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or

(D) for the preparation of a compound of formula (I) in which R^3 represents a hydrogen atom, reacting a compound of formula (VI)

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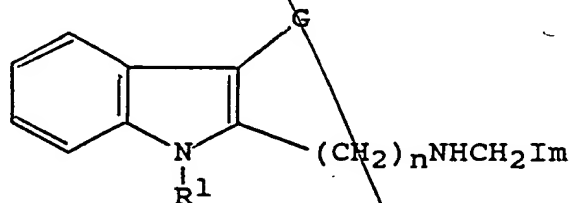
(VI)

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or a protected derivative thereof, with formamide, followed if necessary by removal of any protecting groups present; or

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(E) reacting a compound of formula (VII)



(VII)

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where G represents a hydrogen atom, or a protected derivative thereof, with phosgene in the presence of a Lewis acid; or

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where G represents a bromine or iodine atom, or a protected derivative thereof, with carbon monoxide in the presence of a palladium (II) salt; followed if necessary by removal of any protecting groups present; or

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(F) converting a compound of general formula (I) into another compound of formula (I) using conventional

techniques; or

(G) removing protecting group(s) from a protected form of a compound of formula (I);

and when the compound of formula (I) is obtained as a mixture of enantiomers, optionally resolving the mixture to obtain the desired enantiomer;

and/or where the compound of formula (I) is in the form of a free base, optionally converting the free base into a salt.

14. A pharmaceutical composition for treating a condition caused by a disturbance of "neuronal" 5-HT function which comprises an effective amount to relieve said condition of a compound of formula (I) as defined in claim 1 or a physiologically acceptable salt or solvate thereof together with at least one physiologically acceptable carrier or excipient.

15. A pharmaceutical composition for treating nausea and vomiting and/or for promoting gastric emptying which comprises an effective amount to relieve said nausea and vomiting or to promote said gastric emptying of a compound of formula (I) as defined in claim 1 or a physiologically acceptable salt or solvate thereof together with at least one physiologically acceptable carrier or excipient.

16. A pharmaceutical composition according to claim 14 in a form adapted for oral or parenteral administration.

17. A pharmaceutical composition according to claim 14 wherein the active ingredient is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one or a physiologically acceptable salt or solvate thereof.

18. A pharmaceutical composition according to claim 15

in a form adapted for oral or parenteral administration.

19. A pharmaceutical composition according to claim 15 wherein the active ingredient is 2,3,4,5-tetrahydro-5-methyl-2-[(5-methyl-1H-imidazol-4-yl)methyl]-1H-pyrido[4,3-b]indol-1-one or a physiologically acceptable salt or solvate thereof.

20. A method of treating a condition caused by disturbance of "neuronal" 5-HT function which comprises administering to a patient an effective amount of a compound of formula (I) as defined in claim 1 or a physiologically acceptable salt or solvate thereof to relieve said condition.

21. A method of treating nausea and vomiting and/or promoting gastric emptying which comprises administering to a patient an effective amount of a compound of formula (I) as defined in claim 1 or a physiologically acceptable salt or solvate thereof to relieve said nausea and vomiting and/or promote said gastric emptying.

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